

# ECSA - Cohort of Alcohol-Dependent Patients Admitted for Withdrawal: Suicidal Behaviour Study

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## General

### Identification

Detailed name Cohort of Alcohol-Dependent Patients Admitted for Withdrawal: Suicidal Behaviour Study

Sign or acronym ECSA

CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation CNIL: 25/10/2006

### General Aspects

Medical area Psychology and psychiatry

Pathology (details) Alcoholism, suicidal risk, depression

Health determinants Addictions  
Lifestyle and behavior  
Social and psychosocial factors

Keywords suicide attempt, healthcare users, health events, dependency

### Scientific investigator(s) (Contact)

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Organization Inserm - National Institute of Health and Medical

## Collaborations

## Funding

Funding status Public

Details PHRC (hospital clinical research programme); clinical research delegation (DRC)

## Governance of the database

Sponsor(s) or organisation(s) responsible Inserm - National Institute of Health and Medical Research

Organisation status Public

## Additional contact

## Main features

## Type of database

Type of database Study databases

Study databases (details) Cohort study

Database recruitment is carried out by an intermediary A selection of health institutions and services

Database recruitment is carried out as part of an interventional study No

Additional information regarding sample selection. Inclusion method: Prospective  
Inclusion cut-off date: 01/08/2009  
Number of required subjects: [500-1000]  
Number details: 800

## Database objective

Main objective General aim: To measure the impact of suicidal thoughts, behaviour and death in alcohol dependency.  
Secondary aim: To assess clinical (psychiatric comorbidity), neuropsychological (computer IGT test) and genetic (blood sample) risk factors.

Inclusion criteria Alcohol-dependent, admitted for withdrawal.

## Population type

Age	Adulthood (19 to 24 years) Adulthood (25 to 44 years) Adulthood (45 to 64 years) Elderly (65 to 79 years) Great age (80 years and more)
Population covered	Sick population
Gender	Male Woman
Geography area	Regional
French regions covered by the database	Île-de-France
Detail of the geography area	Multicentric cohort throughout France (5 centres)
<b>Data collection</b>	
<b>Dates</b>	
Date of first collection (YYYY or MM/YYYY)	05/2007
Date of last collection (YYYY or MM/YYYY)	09/2011
<b>Size of the database</b>	
Size of the database (number of individuals)	[500-1000[ individuals
Details of the number of individuals	750
<b>Data</b>	
Database activity	Current data collection
Type of data collected	Clinical data Declarative data Biological data
Clinical data (detail)	Direct physical measures Medical registration
Details of collected clinical data	Clinical examination: At baseline and during follow-up. Examination frequency: 1 (year). Information collected during clinical examination: Depression, dependence, etc. (all psychiatric disorders).

Declarative data (detail)	Face to face interview
Details of collected declarative data	Interview questionnaire: At baseline and during follow-up. Interview frequency: 2 to 3 (years). Information collected during interview: Diagnostic interview for genetic studies (DIGS): semi-structured questionnaire. Other information sheet: During study. Frequency of other information sheet: ONCE AT ANALYSIS COMPLETION (Yearly). Information collected through other information sheet: Vital status. Who completes the other information sheet? Friends or relatives and/or general practitioner.
Biological data (detail)	Type of samples collected: Blood (for DNA)
Presence of a biobank	Yes
Contents of biobank	Plasma DNA
Details of biobank content	Biobank: Blood bank, DNA bank
<b>Procedures</b>	
Quality procedure(s) used	Interviews: online entry; clinical examinations: online entry; biological examinations: online entry; consistency request: after electronic data is entered; management of missing data: refer back to patient; physician reminder for follow-up visits? yes; subject reminder for follow-up visits? yes; patients are informed about the use of their data (written).
Participant monitoring	Yes
Details on monitoring of participants	2 years
Links to administrative sources	No
<b>Promotion and access</b>	
<b>Promotion</b>	
<b>Access</b>	
Terms of data access (charter for data provision, format of data, availability delay)	Can data be used by academic teams? No. Can data be used by industrial teams? No.
Access to aggregated data	Access not yet planned

Access to individual data

No access